CLAIMS

- 1. P2X purinergic receptor antagonist for the treatment of demyelinating and autoimmune diseases, preferably multiple sclerosis, in mammals including man.
- 2. P2X purinergic receptor antagonist for the treatment of demyelinating and autoimmune diseases in accordance with claim 1 characterised because the purinergic receptor is preferably a P2X7 receptor.

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3. P2X purinergic receptor antagonist for the treatment of demyelinating and autoimmune diseases in accordance with claim 1 characterised because the antagonist is a wide spectrum antagonist of P2X receptors or a selective antagonist of a P2X7 receptor, such as o-ATP.

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- 4. P2X purinergic receptor antagonist for the treatment of demyelinating and autoimmune diseases in accordance with claim 1 and 3 characterised because the aforementioned antagonist can be selected from between PPADS, iso-PPADS, Suramin, Evans Blue, NF023, NF279, BBG, NF449, o-ATP, KN62, PPNDS, RB2, MRS2220, Ip51, TNP-ATP or HMA.
- 5. Use of an antagonist of P2X purinergic receptors in the preparation of a drug for the treatment of demyelinating and autoimmune diseases, preferably multiple sclerosis, in mammals including man.

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- Use of an antagonist of P2X purinergic receptors in accordance with claim
 characterised because the aforementioned purinergic receptors are preferably P2X7 receptors.
- 7. Use of an antagonist of P2X purinergic receptors in accordance with claim 5 characterised because the aforementioned antagonist is a wide spectrum antagonist for P2X receptors or a selective antagonist of a P2X7 receptor, such as o-ATP.
- 8. Use of an antagonist of P2X purinergic receptors in accordance with claim

5 to 7 characterised because the aforementioned antagonist can be selected from between PPADS, iso-PPADS, Suramin, Evans Blue, NF023, NF279, BBG, NF449, o-ATP, KN62, PPNDS, RB2, MRS2220, Ip51, TNP-ATP or HMA.

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- 9. A pharmaceutical composition which comprises of at least one P2X purinergic receptor antagonist and at least one pharmaceutically acceptable excipient.
- 10. A pharmaceutical composition in accordance with claim 9 characterised because the antagonist is a wide spectrum antagonist for P2X receptors or a selective antagonist of a P2X7 receptor, such as o-ATP.
- 11. A pharmaceutical composition in accordance with claim 9 to 10 characterised because the aforementioned antagonist is selected from between PPADS, iso-PPADS, Suramin, Evans Blue, NF023, NF279, BBG, NF449, o-ATP, KN62, PPNDS, RB2, MRS2220, Ip51, TNP-ATP or HMA.